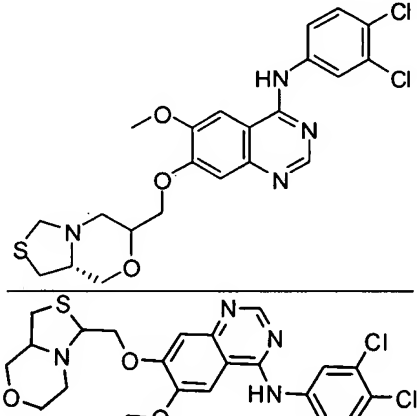


**AMENDMENTS**

**In the Specification**

Please make the following additions to entry 56 in Table 1 on page 23 of the Specification.

56	<p>N-(3,4-dichlorophenyl)-6-(methyloxy)-7-((8aR)-tetrahydro-1H-[1,3]thiazolo[4,3-c][1,4]oxazin-6-ylmethyl)oxy}quinazolin-4-amine;  <u>N-(3,4-dichlorophenyl)-6-(methyloxy)-7-[(tetrahydro-1H-[1,3]thiazolo[4,3-c][1,4]oxazin-3-ylmethyl)oxy]quinazolin-4-amine</u></p>	
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Please amend paragraph [0302] on pages 132-3 of the Specification as follows.

**[0302]** *N*-(3,4-Dichlorophenyl)-7-{[(3-*exo*)-8-methyl-8-azabicyclo[3.2.1]oct-3-yl]oxy}-6-(methyloxy)quinazolin-4-amine hydrochloride: A solution of 4-[(3,4-dichlorophenyl)amino]-6-(methyloxy)quinazolin-7-ol trifluoroacetate (salt) (0.150 g, 0.322 mmol), (3-*endo*)-3-[(methylsulfonyl)methyloxy]-8-azabicyclo[3.2.1]octane (0.106 g, 0.483 mmol), and potassium carbonate (0.220 g, 1.60 mmol) in *N,N*-dimethylacetamide (1.1 mL) was heated in a sealed tube at 100°C for 12 h, followed by 48 h at room temperature. The crude reaction mixture was filtered through celite using methanol eluent, and the solvents were removed *in vacuo*. The residue was purified by HPLC (reverse-phase, acetonitrile/water/0.1% TFA). Upon removal of solvent, the product was taken up in methanol and treated with Bio-Rad AG 1-X8 resin (hydroxide form) until pH 8. The product was filtered and concentrated *in vacuo*, then taken up in methanol and treated with 4.0 M hydrogen chloride in dioxane (0.050 mL). Removal of solvent *in vacuo* provided 48.7 mg (31%) of *N*-(3,4-dichlorophenyl)-7-{[(3-*exo*)-8-methyl-8-azabicyclo[3.2.1]oct-3-yl]oxy}-6-(methyloxy)quinazolin-4-amine hydrochloride. <sup>1</sup>H NMR (400 MHz, d<sub>6</sub>-DMSO): 10.69 (s, 1H), 8.92 (s, 1H), 8.32 (s,

1H), 8.17 (d, 1H), 7.81 (m, 2H), 7.75 (d, 1H), 5.05 (m, 1H), 4.02 (s, 3H), 2.69 (d, 2H),  
2.39 (m, 1H), 2.29-2.18 (m, 6H); MS (EI) for C<sub>23</sub>H<sub>24</sub>N<sub>4</sub>O<sub>2</sub>Cl<sub>2</sub>: 459 (MH<sup>+</sup>).